Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1 (currently amended):

A compound of the formula

wherein

R¹ represents a hydrogen atom or a group selected from the formulae (A) and (B)

(A) R^3 -CO-(CH₂)_s-CO-,

in which

 R^3 represents R^4 – Z^1 with Z^1 being O or NR^5 , R^4 , R^5 being each independently hydrogen or C_{1-6} alkyl, and s is an integer from 1 to 4;

(B) R^6 -CO-

in which

 R^6 represents a C_{1-6} alkyl group, a C_{1-6} haloalkyl group or a phenyl group being optionally substituted by one or more substituents selected from the group consisting of halogen, C_{1-6} alkyl, C_{1-6} alkoxy, C_{1-6} haloalkyl, C_{1-6} haloalkoxy, amino, C_{1-6} alkylamino, di-(C_{1-6} alkyl)-amino, C_{1-6} alkoxycarbonyl, formyl, carboxy, hydroxy, cyano, SO_3H and nitro;

Xaa¹ each independently represent an amino acid or the N-alkylated derivative thereof, at least one of which being N-terminally linked to R¹;

n is 0 or an integer from 1 to 3;

Y represents a single bond, or if t is 0, a spacer group selected from -O- and -NH-; R² represents a hydroxy group or a group of formula (C)

(C) $-Z^2-R^7$

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in which

Z² represents O or NR⁸,

R⁷ represents

- (a) a C₁₋₆ alkyl group being optionally substituted by one or more substituents selected from the group consisting of halogen, C₃₋₈-cycloalkyl, phenyl, C₁₋₆ alkoxy, C₁₋₆ haloalkoxy, amino, C₁₋₆ alkylamino, di-(C₁₋₆ alkyl)-amino, C₁₋₆ alkoxycarbonyl, formyl, carboxy, hydroxy, cyano and nitro, or
- (b) a phenyl group being optionally substituted by one or more substituents selected from the group consisting of halogen, C₁₋₆ alkyl, C₁₋₆ alkoxy, C₁₋₆ haloalkyl, C₁₋₆ haloalkoxy, amino, C₁₋₆ alkylamino, di-(C₁₋₆ alkyl)-amino, C₁₋₆ alkanoylamino, C₁₋₆ alkoxycarbonyl, formyl, carboxy, hydroxy, cyano and nitro,

R⁸ represents a hydrogen atom or C₁₋₆ alkyl group;

Xaa² each independently represent an amino acid or the N-alkylated derivative thereof, in which the amino group of the N-terminally amino acid may have been replaced by Y, and one of which being C-terminally linked to R²;

t is 0 or an integer from 1 to 3;

X is selected from ethyl, thiomethyl and C₃-C₈-cycloalkyl; and

m is 1 or 2,

or a pharmaceutically acceptable salt or solvate thereof.

Claim 2 (original): A compound according to claim 1, wherein Xaa¹ each independently is selected from the group of amino acids consisting of: Leu, Ile, Nva, Abu, Glu, Tle, Phg, Val, allo-Ile, Cpa, Met, Thr, Chg, S-Methylcystein, D-Leu, Nip, CBA (Cyanobutyric acid) and Allyl-Glycin; and n is 1 or 2.

Claim 3 (original): A compound according to claim 1, wherein Xaa² each independently is selected from the group of amino acids consisting of: Val, Ala, Leu, Ile, Nva, Abu, Cha, Tle, Phg, Glu, Nle, Phe, His, Ser, Cpa, and Asp; and s is 1 or 2.

Claim 4 (original): A compound according to claim 2, wherein

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Xaa² each independently is selected from the group of amino acids consisting of: Val, Ala, Leu, Ile, Nva, Abu, Cha, Tle, Phg, Glu, Nle, Phe, His, Ser, Cpa, and Asp; and s is 1 or 2.

Claim 5 (original): A compound according to claim 1, wherein m represents 1.

Claim 6 (original): A compound selected from the formulae (IA) through (ID):

$$R^{1}-(Xaa^{1})_{\overline{n-1}}N = C \qquad \qquad N \qquad \qquad N \qquad \qquad C \qquad \qquad (IC)$$

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in which R¹, R², Xaa¹, Xaa², n and t are as defined in claim 1, and X represents ethyl, thiomethyl or cyclopropyl; or a pharmaceutically acceptable salt or solvate thereof.

Claim 7 (original): A pharmaceutical composition comprising a compound according to claim 1 or a pharmaceutically acceptable salt or solvate thereof; and a pharmaceutically acceptable carrier or diluent.

Claim 8 (original): A pharmaceutical composition comprising a compound according to claim 6 or a pharmaceutically acceptable salt or solvate thereof; and a pharmaceutically acceptable carrier or diluent.

Claim 9 (original): A pharmaceutical composition according to claim 7, which further comprises an active ingredient selected from the group consisting of: atorvastatin, besipirdine, cevimeline, donepezil, eptastigmine, galantamine, glatiramer acetate, icopezil, ipidacrine, lazabemide, linopirdine, lubeluzole, memantine, metrifonate, milameline, nefiracetam, nimodipine, octreotide, rasagiline, rivastigmine, sabcomeline, sabeluzole, tacrine, valproate sodium, velnacrine, YM 796, Phenserine and zanapezil.

Claim 10 (currently amended): A pharmaceutical composition according to claim 7, which further comprises an antiinflammtory agent selected from the group consisting of: rofecoxib, celecoxib, valdecoxib, nitroflurbiprofen, IQ-201, NCX-2216, CPI-1189, a complex of proline-rich polypeptides derived from ovine colostrums and sold under the trademark Colostrinin, ibuprofen, indomethacin, meloxicam, and sulindac sulphide.

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Claim 11 (currently amended) A pharmaceutical composition according to claim 9, which further comprises an antiinflammtory agent selected from the group consisting of: rofecoxib, celecoxib, valdecoxib, nitroflurbiprofen, IQ-201, NCX-2216, CPI-1189, a complex of proline-rich polypeptides derived from ovine colostrums and sold under the trademark Colostrinin, ibuprofen, indomethacin, meloxicam, and sulindac sulphide.

Claim 12 (original): A pharmaceutical composition according to claim 7, which further comprises a nerve growth factor or a nerve growth modulator selected from the group consisting of: ABS-205, Inosine, KP-447, leteprinim, MCC-257, NS-521, and xaliproden.

Claim 13 (original): A pharmaceutical composition according to claim 9, which further comprises a nerve growth factor or a nerve growth modulator selected from the group consisting of: ABS-205, Inosine, KP-447, leteprinim, MCC-257, NS-521, and xaliproden.

Claim 14 (original): A pharmaceutical composition according to claim 11, which further comprises a nerve growth factor or nerve growth modulator selected from the group consisting of: ABS-205, Inosine, KP-447, leteprinim, MCC-257, NS-521, and xaliproden.

Claims 15-18 (canceled)